FDA 510(k) Premarket Notification

K832827 CORPULS 300

CLEARED

Nov 3, 1983
73 days to decision

K832827 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k832827/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 22, 1983
Decision date	Nov 3, 1983
Days to decision	73 days
Third-party review	No

APPLICANT

Company	**Automated Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1983